## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

## Listing of Claims:

1. (Original) A compound of formula I, pharmaceutically acceptable salts thereof, or mixtures thereof:

wherein

 $R^1$  is an aryl, heteroaryl, substituted aryl or substituted heteroaryl; and  $R^2$  is hydrogen, optionally substituted  $C_{1-12}$ alkyl, optionally substituted  $C_{6-12}$ aryl, or optionally substituted  $C_{2-12}$ heterocyclyl.

2. (Original) A compound according to claim 1,

wherein  $R^1$  is selected from phenyl; pyridyl; thienyl; furyl; imidazolyl; triazolyl; pyrrolyl; thiazolyl; and N-oxido-pyridyl, optionally substituted with one or more groups selected from  $C_{1-6}$ alkyl, halogenated  $C_{1-6}$ alkyl, -NO<sub>2</sub>, -CF<sub>3</sub>,  $C_{1-6}$  alkoxy, chloro, fluoro, bromo, and iodo; and

R<sup>2</sup> is hydrogen or methyl.

3. (Original) A compound according to claim 1,

wherein  $R^1$  is selected from phenyl; pyridyl; thienyl; furyl; imidazolyl; pyrrolyl; and thiazolyl, optionally substituted with one or more groups selected from  $C_{1-6}$ alkyl, halogenated  $C_{1-6}$ alkyl, -NO<sub>2</sub>, -CF<sub>3</sub>,  $C_{1-6}$  alkoxy, chloro, fluoro, bromo, and iodo; and

R<sup>2</sup> is hydrogen or methyl.

4. (Original) A compound according to claim 1,
wherein R<sup>1</sup> is selected from phenyl; pyridyl; thienyl; furyl; imidazolyl;
pyrrolyl; and thiazolyl; and
R<sup>2</sup> is hydrogen or methyl.

5. (Original) A compound according to claim 1, wherein the compound is selected from:

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3-[(4-[(diethylamino)carbonyl]phenyl)(4-benzyl-piperazin-1-yl)methyl]benzamide;
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- 3-{(4-[(diethylamino)carbonyl]phenyl)[4-(2-furylmethyl)-piperazin-1-yl]methyl}benzamide;
- 3-[[4-[(diethylamino)carbonyl]phenyl][4-(phenylmethyl)-1-piperazinyl]methyl]-N-methyl-benzamide; enantiomers thereof; and pharmaceutically acceptable salts thereof.
- 6-7. (Cancelled)
- 8. (Currently Amended) A pharmaceutical composition comprising a compound according to any one of claims 1–5claim 1 and a pharmaceutically acceptable carrier.
- 9. (Currently Amended) A method for the therapy of pain in a warm-blooded animal, comprising the step of administering to said animal in need of such therapy a therapeutically effective amount of a compound according to any one of claims 1-5claim 1.
- 10. (Currently Amended) A method for the therapy of functional gastrointestinal disorders in a warm-blooded animal, comprising the step of administering to said

animal in need of such therapy a therapeutically effective amount of a compound according to any one of claims 1-5claim 1.

## 11. (Original) A process for preparing a compound of formula II,

comprising of the step of reacting a compound of formula III:

 $\overline{\mathbf{III}}$ 

with R<sup>1</sup>-CHO to form the compound of formula II wherein

R<sup>1</sup> is an aryl, heteroaryl, substituted aryl or substituted heteroaryl.

## 12. (Original) A process for preparing a compound of formula IV,

comprising: reacting a compound of formula II,

with an akali metal hydroxide in non-aqueous solvent to form the compound of formula IV:

wherein

 $R^{1}$  is an aryl, heteroaryl, substituted aryl or substituted heteroaryl.